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| <p>(54) Title: DIALYSATE COLLECTION BAG AND METHOD OF STERILIZING FLUID COLLECTED THEREIN</p> <p>(57) Abstract</p> <p>A method of peritoneal dialysis which utilizes a dialysis collection bag (10) connected to a drain tube (12) is disclosed. The collection bag (10) is sealed so that it is liquid tight and liquid cannot enter or leave it except through drain tube (12). Inside the collection bag (10) is a sterilization tablet (14) which preferably contains calcium hypochlorite of sufficient quantity to quickly sterilize 2 liters to 4 liters of used dialysis solution.</p> <div data-bbox="971 1155 1412 1852"><p>The diagram shows a rectangular dialysis collection bag (10) with a drain tube (12) extending from the top. At the bottom of the bag, there is a small, rounded compartment containing a sterilization tablet (14).</p></div> | | |

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**DIALYSATE COLLECTION BAG AND
METHOD OF STERILIZING FLUID COLLECTED THEREIN**

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FIELD OF THE INVENTION

10 The present invention relates to the field of
medical dialysis including peritoneal dialysis and
hemodialysis. In particular, the present invention
relates to a dissolving bacteriocidal additive to a fluid
collection bag to sterilize the fluid collected in the
15 bag for safe disposal of the sterilized fluid.

BACKGROUND OF THE INVENTION

 The common treatment for renal failure is
hemodialysis treatment or peritoneal dialysis treatment.
20 Both treatments utilize the diffusion of liquid through a
semipermeable membrane. In the case of hemodialysis the
membrane is in a dialyzer external to the patient, so
that blood is withdrawn from the patient's vascular
system and passed across the membrane while dialysis
25 solution is passed across the other side of the membrane.
Impurities in the blood are drawn through the membrane by
osmotic pressure on the membrane and are disposed of in
the discarded dialysis solution. In the case of
peritoneal dialysis, the semipermeable membrane is the
30 patient's peritoneal membrane. Dialysis solution is
introduced into and retained for a period of time in the
peritoneal cavity, and impurities in the blood migrate
through the peritoneal membrane and into the dialysis
solution. The dialysis solution with the impurities is
35 then withdrawn from the peritoneal cavity and discarded
into a "drain" bag or "collection" bag.

 Both hemodialysis and peritoneal dialysis
require significant amounts of dialysis solution,

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sometimes called dialysate. Common dialysates are primarily water, but with low ionic concentrations of dissolved sodium, potassium, calcium, magnesium, chloride, acetate, glucose and bicarbonate. The proportions of these and other compounds depends on a variety of factors. Regardless of the exact concentrations and relative proportions of the dissolved compounds, the main material in all dialysates is water.

Dialysis solutions have been premixed and prepackaged in a variety of mixes and sizes, so that the patient or the medical professional simply selects the desired size and mix of dialysate, makes the appropriate tubing connections to the prepackaged dialysates and to the patient and the dialysis machine, and then commences the procedure.

Regardless of the type of dialysis - peritoneal dialysis or hemodialysis - the procedure results in significant and sometimes substantial quantities of spent dialysis solution. This used dialysis solution has been exposed to and often contains a variety of pathogens including infectious diseases. There are cases in which used dialysis solution was found to contain viable HIV virus, for example, and there are undoubtedly other instances where used dialysis solution contained bacteria or viruses of other diseases as well.

This used dialysis solution is normally deemed "medical waste" and is required to be disposed of in accordance with applicable medical waste disposal procedures so as not to spread disease or contaminate water supplies. However, it is commonly recognized that the used dialysis solution is often disposed of improperly. Because many dialysis procedures are adapted for the convenience of home use, used dialysis solution is sometimes improperly discarded by simply putting it into trash receptacles for ordinary trash pick-up or

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pouring it into a sink or flushing it down the toilet. Even in hospitals and clinics, the high cost and inconvenience of medical waste disposal may tempt professionals to dispose of used dialysis solution improperly.

In the field of urinary catheters, there is a body of art pertaining to preventing pathogens from migrating from a urine collection bag up through a catheter and into the urethra. Such art is not directed toward the sterilization of the collected urine so that it can be properly disposed of without infecting others since urine does not normally contain infectious diseases even if from a diseased patient. Instead, this art is more directed toward preventing the collection bag from becoming a colonization site from which infection can migrate up the catheter to the patient himself. Illustrative of this urinary catheter art are U.S. Patent Nos. 4,529,398 by Wong; 4,661,100 by Rechsteiner; 5,267,989 by Moyet-Ortiz; 4,863,445 by Mayhan; 4,417,892 by Meisch; and 4,372,313 by Villari.

The typical approach to preventing urinary tract infections in the urinary catheter prior art mentioned above is to include a sterilizing agent in the catheter or in the collection bag so that pathogens cannot migrate up the catheter. Such an approach is not appropriate for a dialysis collection bag, however, because in dialysis it is often desired to take specimen samples of the used dialysate. If the dialysate is sterilized upon entering the collection bag, then specimen samples cannot be cultured to test for live pathogens. It is also desirable in dialysis that the collection bag be entirely self-contained, so that the collection bag is manufactured with the sterilizing agent pre-placed within it to avoid a separate placement step at the time of use.

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In the Wong patent, a dispensing device having a polymer with a chemoprophylactic agent is placed within the collection bag. The dispensing device begins sterilizing liquid in the collection bag immediately upon contact, and the device is designed such that the sterilizing properties continue for an extended period of time. In contrast, in dialysis collection bags it is desirable that the sterilization of the liquid not be commenced immediately upon contact with the dialysis collection bag, and in dialysis collection bags it is not necessary that the sterilization be contained for an extended period of time because the bag is filled in a very short period of time rather than over a period of many hours.

The Rechsteiner patent discloses a system with a urine collection bag having a fragile resinous material inside which is broken to release a sterilizing or diagnostic agent. The Rechsteiner patent is like the Wong patent in that it is designed for urinary catheter applications in which the collected urine must be sterilized immediately upon contact and over an extended period of time to prevent pathogen migration into the patient. The Mayhan patent is similar to the Rechsteiner patent, except that the resinous sterilizing agent is replaced with a slow-dissolving tablet. The Moyet-Ortiz patent discloses an antiseptic absorbent pad in a urine collection device; the Meisch patent discloses an outlet tube to a urine collection bag which is treated with a sterilizing agent to prevent pathogen colonization; the Villari patent discloses a urine collection bag with a tubular portion having a device for retaining an antimicrobial agent.

SUMMARY OF THE INVENTION

The present invention includes a dialysis

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collection bag and method of using the same in dialysis treatment and the disposal of used dialysis fluid. The apparatus of the invention includes a used dialysate collection bag having a sterilizing tablet within the collection bag. The sterilizing pill is coated with a gelatinous coating to delay the dissolution of the pill when the collection bag is filled.

In an alternative embodiment, the present invention includes a used dialysate collection bag and a sterilization bag connected thereto. The sterilization bag contains a sterilizing agent and is attached to the collection bag by a breakable connection. Once the connection is broken, there is fluid communication between the collection bag and the sterilization bag.

The system is operated by performing peritoneal dialysis in the conventional manner, including the step of draining the used dialysis solution into the collection bag. In one embodiment, the used dialysis solution begins to dissolve the gelatinous coating on the sterilization tablet immediately upon entering the collection bag. However, the gelatinous coating on the sterilization tablet is such that the tablet does not release the sterilization agent for about 30 minutes after the initial contact between the used dialysis solution and the sterilization tablet. This 30 minute delay allows time for a specimen sample to be taken from the collection bag before the sterilization agent is released to sterilize the used dialysis solution. Once the approximately 30 minutes expires so that the gelatinous coating is dissolved and the sterilization agent is released from the sterilization tablet, the sterilization agent is completely released, substantially immediately. In an alternative embodiment, when the sterilization agent is contained in a sterilization bag attached to the collection bag by a breakable connector,

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once the collection bag has been filled and samples taken the connection between the two bags is broken and the sterilization agent is completely released into the dialysis solution. In either embodiment, unlike in the case of urine collection bags used with urinary catheters, there is no need for slow and prolonged release over a period of time. The collection bag with sterilized used dialysis solution is disposed of in the standard proper manner as medical waste, but in the event that the disposal is improper for some reason, there is now much less likelihood of spreading infectious disease.

A variety of sterilization agents and gelatinous coatings are feasible for the sterilization tablet. In a preferred embodiment, the sterilization agent is calcium hypochlorite and the coating is sodium laurel sulfate.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 shows a perspective view of a dialysis collection bag in accordance with the present invention

FIG. 2 shows a perspective, partial cut-away view of a sterilization tablet in accordance with the present invention.

FIG. 3 shows a perspective view of an alternative embodiment of the invention, showing a sterilization bag attached to the dialysis collection bag by a frangible seal.

DETAILED DESCRIPTION OF THE INVENTION

A perspective view of a dialysis collection bag in accordance with the present invention is shown in FIG. 1. A dialysis collection bag 10 is a bag-shaped element connected to a drain tube 12 which is connected to an ordinary peritoneal dialysis or hemodialysis tubing set (not shown). The collection bag 10 is sealed so that it

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is liquid tight and liquid cannot enter or leave it except through the drain tube 12. The collection bag 10 is constructed from plastic sheet material in the conventional manner known in the art and is sized appropriately so that the collection bag alone or in combination with a collection bag set has sufficient capacity to receive the used dialysis solution from a dialysis treatment.

Inside the collection bag 10 is a sterilization tablet 14. The sterilization tablet 14 is preferably placed within the collection bag 10 at the time the collection bag is being manufactured. The sterilization tablet 14 is placed in the collection bag 10, and then the collection bag 10 is sealed to retain the sterilization tablet 14 therein. In a preferred embodiment, the tablet contains calcium hypochlorite. Other disinfectant agents are feasible as well, including lithium hypochlorite, sodium hypochlorite, and powder chloramines such as sodium p-toluene sulfonchloramide or other soluble disinfectants. The quantity and concentration of disinfectant should be sufficient to quickly disinfect 2-4 liters of used dialysis solution contained within the collection bag.

An important aspect of the preferred embodiment of the invention is the use of a dissolving coating 16 over the body 18 of the sterilization tablet 14 as shown in FIG. 2. The coating must be dissolved by contact with the used dialysis solution before the disinfectant in the sterilization tablet 14 is released. As explained above, this delays the activation of the sterilization tablet 14 for a short time after initial contact with the used dialysis solution, to allow time to take a specimen sample before the disinfecting process commences. It has been found that the most convenient time period to provide between the initial contact between the

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sterilization tablet 14 and the used dialysis solution, to allow for the collection of a specimen sample, is about 30 minutes.

5 The coating 16 in the preferred embodiment is sodium laurel sulfate or gelatin. Other coatings are also feasible to provide appropriate delay in activation of the sterilization tablet 14, including ethylcellulose, hydroxypropyl-methylcellulose, titanium dioxide, sucrose stearate, hydroxypropylcellulose, polyvinyl-
10 acetal-diethylaminoacetate, or acrylic latex sprays. The coating should dissolve in a pH range of 3-10.

 The drain tube 12, as well as the rest of the tubing set and the collection bag 10 as well, may be
15 sterilized at the time of manufacture by processes known in the art. Such processes include but are not limited to the use of ethylene oxide gas, steam, gamma rays and electron beams.

 The manufacturing method will normally include the fabrication and assembly of the tubing set with its
20 various elements such as valves, clamps and so on; the construction of the collection bag 10 with the sterilization tablet 14 therein; the attachment of the collection bag 10 to the drain tube 12; and the sterilization of the entire assembly by one or more of
25 the known means mentioned above. The assembly is used in peritoneal dialysis and hemodialysis procedures much like tubing sets of the prior art. When the collection bag 10 is filled with used dialysis solution, the coating 16 to the sterilization tablet 14 begins to dissolve. The time
30 it takes for the coating 16 to dissolve to the point where the sterilization agent is released may vary, as mentioned above, from a few seconds to an hour or longer. Preferably, the time is at least a few minutes, in order to allow time for a specimen sample to be taken, and the
35 time is not so long that the used dialysis solution may

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be removed from the collection bag 10 prior to being sterilized by release of the sterilization agent. It has been found that about 30 minutes is ideal for this time period.

5 Once any desired specimen sample has been taken, and the coating 16 to the sterilization tablet 14 is dissolved so that the sterilization agent is released from the sterilization tablet 14 to sterilize the used dialysis solution, the collection bag 10 with the used,
10 sterilized dialysis solution therein is discarded. It is still desirable that the collection bag 10 with the used, sterilized dialysis solution therein be discarded as medical waste, in accordance with proper medical waste disposal procedures. However, the prior sterilization of
15 the used dialysis solution provides some assurance against the spread of disease in the event that such procedures are ignored or followed improperly.

 With reference now to FIG. 3, an alternative embodiment of the present invention is readily
20 understood. In this embodiment, a sterilization bag 20 is attached to the collection bag 10 by a frangible seal 22 which separates the bags. Preferably, the sterilization agent will be placed with the sterilization bag 20 when the sterilization bag and the collection bag
25 are being manufactured. As mentioned previously, the quality and concentration of disinfectant contained within the sterilization bag 20 should be sufficient to quickly disinfect 2-4 liters of used dialysis solution contained within the collection bag 10. A sealing device,
30 preferably of the heating type, will be employed to seal the walls of frangible seal 22 so that the sterilization agent is separate from the initially empty collection bag 10. The frangible seal 22 is readily delaminated to permit the free flow of materials between the containers
35 while securing the containers to each other.

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As described herein, peritoneal dialysis is performed on a patient and the used dialysis solution is collected in the collection bag 10. While the dialysis solution is collected and the collection bag 10 is filled, any desired specimen samples may be taken. Once the desired specimen samples have been taken, the frangible seal 22 between the collection bag 10 and the sterilization bag 20 is broken, providing free flow of materials between the compartments. The used dialysis solution is sterilized substantially immediately. The sterilization bag 20 and the collection bag 10 with the used dialysis solution therein are discarded.

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CLAIMS

What is claimed is:

1. A method of peritoneal dialysis,
comprising: engaging with a dialysis patient a dialysis
5 tubing set including a dialysis solution source and a
collection bag having a sterilization device therein;
performing the dialysis on the patient using the dialysis
solution; draining the used dialysis solution into the
collection bag; and sterilizing the used dialysis
10 solution with the sterilization device in the collection
bag.
2. The method of claim 1, wherein the step of
sterilizing the used dialysis solution is by a release of
a sterilization agent from the sterilization device a
15 known period of time after contact between the
sterilization device and the used dialysis solution.
3. The method of claim 2, wherein the
sterilization device includes a sterilization tablet with
a coating, and wherein the sterilization step includes
20 dissolving the coating with the used dialysis solution
before releasing the sterilization agent.
4. The method of claim 3, wherein the
sterilization agent is a soluble disinfectant.
5. The method of claim 4, wherein the
25 sterilization agent is selected from the group consisting
of calcium hypochlorite, lithium hypochlorite, sodium
hypochlorite and powder chloramines.
6. The method of claim 4, wherein the coating
is selected from the group consisting of sodium laurel
30 sulfate, gelatin, ethylcellulose,
hydroxypropylmethylcellulose, titanium dioxide, sucrose
stearate, hydroxypropylcellulose, polyvinyl-
acetal-diethylaminoacetate or acrylic.
7. The method of claim 4, wherein the coating
35 takes at least thirty minutes to dissolve.

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8. A method of performing peritoneal dialysis on a patient, using a peritoneal dialysis solution, and safely disposing of the peritoneal dialysis solution, comprising: performing peritoneal dialysis on the patient
5 using the peritoneal dialysis solution; draining the peritoneal dialysis solution into a collection bag having a sterilizing compound therein for sterilizing the used dialysis solution; sterilizing all the used peritoneal dialysis solution drained into the collection bag by
10 contact of the peritoneal dialysis solution with the sterilizing compound; and discarding the sterilized peritoneal dialysis solution.

9. A method of peritoneal dialysis, comprising: engaging with a dialysis patient a dialysis
15 tubing set including a dialysis solution source and a collection bag having a sterilization bag attached by a frangible seal thereto, the sterilization bag containing a sterilization agent therein; performing the dialysis on the patient using the dialysis solution; draining the
20 used dialysis solution into the collection bag; breaking the seal between the collection bag and the sterilization bag causing fluid communication therebetween; and sterilizing the used dialysis solution with the sterilization agent in the sterilization bag.

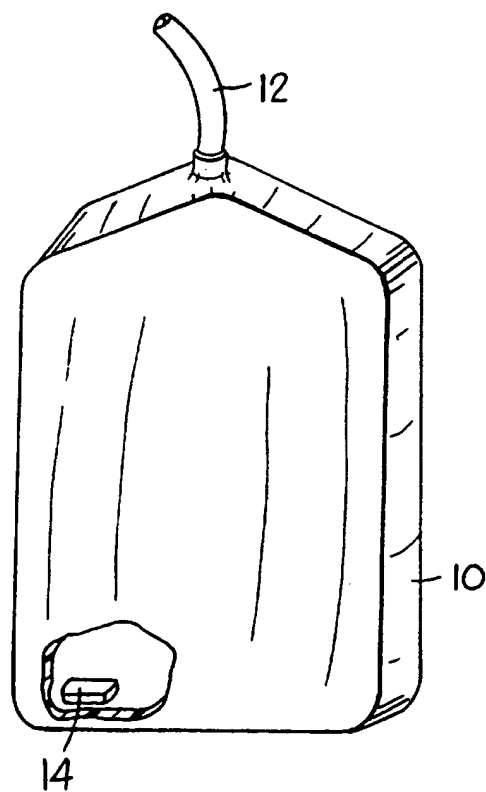


FIG. 1

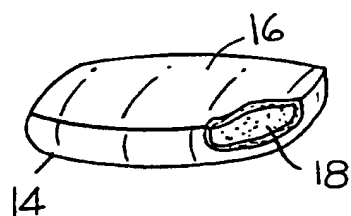


FIG. 2

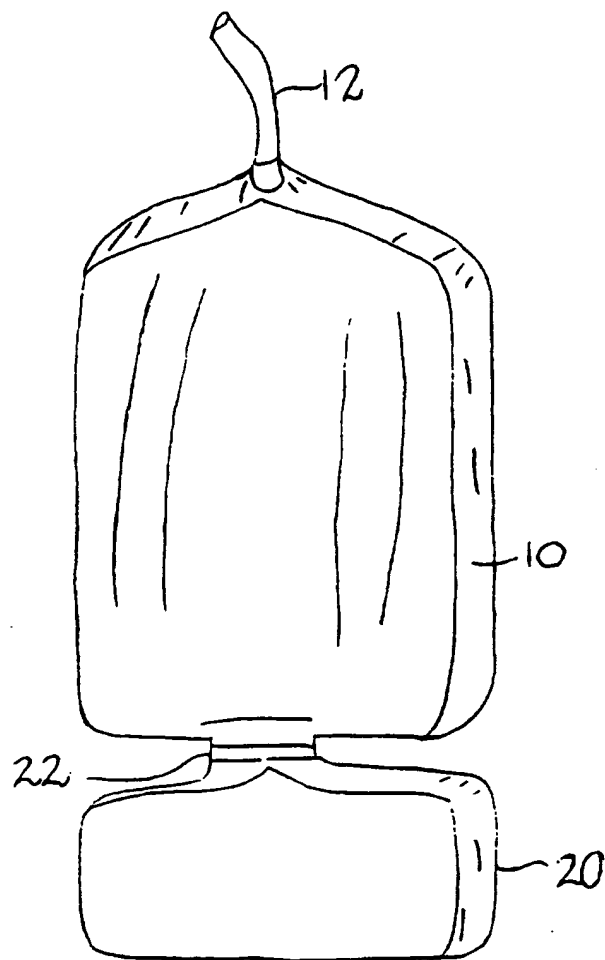


FIG. 3

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US96/15534

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :A61M 37/00

US CL :604/4

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 128/DIG. 24; 604/4, 28, 29, 416

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

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C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category* | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|-----------|--|-----------------------|
| Y | US 4,190,047 A (JACOBSEN et al) 26 February 1980, entire document. | 1, 2, 8 |
| Y | US 4,529,398 A (WONG et al) 16 July 1985, entire document. | 1, 2, 8 |

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

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Facsimile No. (703) 305-3590

Authorized officer

PAUL PREBILIC

Telephone No. (703) 308-2905